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39277 casein and nanoparticle\$2 DERMENT	7 13 19 25	35277 401 35 10676 591	and nanoparticle\$2 and nanoparticle\$2.clm. same gelatin same gelatin.clm.	USPAT; USPAT; US-PGPUB; EPO; JPO; DERWENT US-PGPUB; EPO; JPO; DERWENT USPAT; US-PGPUB;	13 14 14
401 casein and nanoparticle\$2	13 19 25	401 35 36 10676 591	and nanoparticle\$2 and nanoparticle\$2.clm. same gelatin same gelatin.clm.	USPAT;	14 14 13
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401 casein and nanoparticle\$2 DERWENT	13 19 25	401 35 10676 591	and nanoparticle\$2 and nanoparticle\$2.clm. same gelatin same gelatin.clm.	DERWENT USPAT; US-PGPUB; EPO, JPO; DERWENT USPAT; US-PGPUB; EPO; JPO; DERWENT USPAT; US-PGPUB; EPO; JPO; DERWENT USPAT; US-PGPUB; EPO; JPO; DERWENT USPAT; US-PGPUB;	13
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35 Gasein and nanoparticle\$2.clm. USPAT; 10676 Casein same gelatin.clm. USPAT; 10676 USPAT; USPAT; 10677 USPAT; USPAT; 10678 USPAT; USPAT; 10679 USPAT; USPAT; USPAT; 10679 USPAT; USPAT; USPAT; USPAT; 10679 USPAT; U	25	35 10676 591 15	and nanoparticle\$2.clm. same gelatin same gelatin.clm.	USPAT; US-PGPUB; EPO; JPO; DERWENT USPAT; US-PGPUB; EPO; JPO; DERWENT USPAT; USPAT; US-PGPUB;	
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See		591	gelatin.clm.	US-PGPUB; EPO; JPO; DERWENT USPAT; US-PGPUB; EPO; JPO;	
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403 Breitenbach.in. USPAT; USPAT; US-PGPUB; EPO; JPO; DERWENT USPAT; US-PGPUB; US-PGPUB; US-PGPUB; US-PGPUB; US-PGPUB; US-PGPUB; EPO; JPO; DERWENT US-PGPUB; EPO; JPO; DERWENT			<u> </u>	EPO; JPO;	
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1 Breitenbach.in. and nanoparticle\$2 USPAT; US-PGPUB; EPO; JPO; US-PGPUB; EPO; JPO; DERWENT				US-PGPUB;	
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1	345	nanoparticle\$2 and polymer\$2.clm. and gelatin	USPAT;	2003/04/02 11:02
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			DERWENT	
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			EPO; JPO;	
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NEWS 44

Feb 24

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METADEX enhancements

PCTGEN now available on STN

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NEWS 45 Feb 24 TEMA now available on STN NEWS 46 Feb, 26 NTIS now allows simultaneous left and right truncation NEWS 47 Feb 26 PCTFULL now contains images NEWS 48 Mar 04 SDI PACKAGE for monthly delivery of multifile SDI results NEWS 49 Mar 19 APOLLIT offering free connect time in April 2003 NEWS 50 Mar 20 EVENTLINE will be removed from STN NEWS 51 Mar 24 PATDPAFULL now available on STN NEWS 52 Mar 24 Additional information for trade-named substances without structures available in REGISTRY Indexing from 1957 to 1966 added to records in CA/CAPLUS NEWS 53 Mar 24 NEWS EXPRESS January 6 CURRENT WINDOWS VERSION IS V6.01a, CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP), AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002 STN Operating Hours Plus Help Desk Availability NEWS HOURS General Internet Information NEWS INTER NEWS LOGIN Welcome Banner and News Items Direct Dial and Telecommunication Network Access to STN NEWS PHONE NEWS WWW CAS World Wide Web Site (general information) Enter NEWS followed by the item number or name to see news on that specific topic. All use of STN is subject to the provisions of the STN Customer agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties. * * * * * * * * * STN Columbus FILE 'HOME' ENTERED AT 15:10:16 ON 03 APR 2003 => file caplus uspatfull biosis europatfull COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 0.21 0.21 FILE 'CAPLUS' ENTERED AT 15:10:53 ON 03 APR 2003 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS) FILE 'USPATFULL' ENTERED AT 15:10:53 ON 03 APR 2003 CA INDEXING COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS) FILE 'BIOSIS' ENTERED AT 15:10:53 ON 03 APR 2003 COPYRIGHT (C) 2003 BIOLOGICAL ABSTRACTS INC. (R) FILE 'EUROPATFULL' ENTERED AT 15:10:53 ON 03 APR 2003 COPYRIGHT (c) 2003 WILA Verlag Muenchen (WILA) => => s nanoparticle? and (gelatin or casein) 1648 NANOPARTICLE? AND (GELATIN OR CASEIN) => s l1 and matrix? 981 L1 AND MATRIX?

=> s 12 and (pharmaceutical or drug? or active agent or peptide?)

=> s 13 and gelatin

L4 896 L3 AND GELATIN

=> s 14 and hydrosol

L5 12 L4 AND HYDROSOL

=> d 15 1-12 abs bib

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ANSWER 1 OF 12 CAPLUS COPYRIGHT 2003 ACS
    Nanoparticulate prepns. of pharmaceutical and cosmetic active
AΒ
     substances with a core-shell structure are described, wherein the active
     substance is present in the core as x-ray amorphous particles dispersed in
     a polymer matrix, and the shell consists of a stabilizing
     sheathing matrix of a swellable polymer. The polymer
     matrix in the core prevents crystn. and pptn. of the active
     substance in the presence of solvent. The polymeric shell maintains the
     core-shell particles in a colloidal state, preventing aggregation or
     flocculation. Thus, cyclosporin A 3 was suspended in a soln. of ascorbyl
     palmitate 0.6 and Kollicoat MAE 0.6 in iso-PrOH 36 g. This suspension was
     then mixed with 120 g H2O at 200.degree. for 0.3 s, followed by mixing
     with a soln. of gelatin A 4.3 and lactose 6.5 in demineralized
     water 490 g (pH \bar{9}.0) at 25.degree. and 30 bar. The resulting dispersion
     (mean particle size 249 nm) was spray dried to a powder (cyclosporin
     content 20.03 wt.%) which could be redispersed in water to a
     hydrosol whose particle size remained stable at 263 nm over 1 h.
AN
     2000:401632 CAPLUS
DN
     133:48938
    Nanoparticulate core-shell systems and use thereof in
ΤI
     pharmaceutical and cosmetic preparations
     Heger, Robert; Auweter, Helmut; Breitenbach, Joerg; Bohn, Heribert
IN
     BASF Aktiengesellschaft, Germany
PA
SO
     PCT Int. Appl., 29 pp.
     CODEN: PIXXD2
DT
     Patent
LΑ
     German
FAN.CNT 1
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     PATENT NO.
                      KIND
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PI
    WO 2000033820
                      A2
                            20000615
                                           WO 1999-EP9545
                                                             19991207
     WO 2000033820
                      A3
                            20001012
         W:
            CA, CN, JP, US
         RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
             PT, SE
     DE 19856432
                            20000615
                                           DE 1998-19856432 19981208
                       Α1
     EP 1137404
                       A2
                            20011004
                                           EP 1999-963399
                                                            19991207
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
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IE, FI

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19981208

19991207

JP 2000-586313

19991207

JP 2002531492

WO 1999-EP9545

PRAI DE 1998-19856432

L5 ANSWER 2 OF 12 USPATFULL

The present invention relates to compounds that specifically block the binding between a member of the HuR family of proteins and a mRNA encoding a member of the CD83 family of proteins and that reduce expression of a member of the CD83 family of proteins in a cell as well as **pharmaceutical** compositions comprising such compounds and methods for screening and/or identifying compounds that block the binding between a member of the HuR family of proteins and a mRNA encoding a member of the CD83 family of proteins.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AN 2002:295140 USPATFULL

TI Compounds that affect CD83 expression, **pharmaceutical** compositions comprising said compounds and methods for identifying said compounds

IN Hauber, Joachim, Langensendelbach, GERMANY, FEDERAL REPUBLIC OF Prechtel, Alexander Thorsten, Lauf an der Pegnitz, GERMANY, FEDERAL REPUBLIC OF

PI US 2002165186 A1 20021107

AI US 2001-25367 A1 20011219 (10)

PRAI GB 2000-31145 20001220

DT Utility

FS APPLICATION

LREP Leopold Presser, Esq., SCULLY, SCOTT, MURPHY & PRESSER, 400 Garden City Plaza, Garden City, NY, 11530

CLMN Number of Claims: 31 ECL Exemplary Claim: 1 DRWN 15 Drawing Page(s)

LN.CNT 2667

ANSWER 3 OF 12 USPATFULL L5

New biologically active compounds are described which inhibit the ΑB cellular formation of niacinamide mononucleotide, and essential intermediate of the NAD(P) biosynthesis in the cell. These compounds can represent the active ingredient of a pharmaceutical composition for the treatment of cancers, leukaemias or for immunosuppression. Furthermore, screening methods are described as a tool for detecting the above active compounds, and for examination of a given cell type for its dependency on niacinamide as a precursor for NAD synthesis.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

2002:288098 USPATFULL AN

ΤI Inhibitors of cellular niacinamide mononucleotide formation and their use in cancer therapy

Biedermann, Elfi, Vaterstetten, GERMANY, FEDERAL REPUBLIC OF IN Eisenburger, Rolf, Kirchseeon, GERMANY, FEDERAL REPUBLIC OF Hasmann, Max, Neuried, GERMANY, FEDERAL REPUBLIC OF Loser, Roland, Feldafing, GERMANY, FEDERAL REPUBLIC OF Rattel, Benno, Munich, GERMANY, FEDERAL REPUBLIC OF Reiter, Friedemann, Putzbrunn, GERMANY, FEDERAL REPUBLIC OF Schein, Barbara, Neufahrn, GERMANY, FEDERAL REPUBLIC OF Schemainda, Isabel, Munich, GERMANY, FEDERAL REPUBLIC OF Schulz, Michael, Aschheim, GERMANY, FEDERAL REPUBLIC OF Seibel, Klaus, Grafelfing, GERMANY, FEDERAL REPUBLIC OF Vogt, Klaus, Munich, GERMANY, FEDERAL REPUBLIC OF Wosikowski, Katja, Poing, GERMANY, FEDERAL REPUBLIC OF

Klinge Pharma GmbH (non-U.S. corporation)

ΡI US 2002160968

A1 20021031

US 6506572

B2 20030114

ΑI US 2001-935772

A1 20010823 (9) Continuation of Ser. No. WO 2000-EP1628, filed on 28 Feb 2000, UNKNOWN

EP 1999-103814 19990226 PRAI

DTUtility

PΑ

RLT

APPLICATION FS ·

FITCH EVEN TABIN AND FLANNERY, 120 SOUTH LA SALLE STREET, SUITE 1600, LREP CHICAGO, IL, 60603-3406

CLMN Number of Claims: 25

Exemplary Claim: 1 ECL

DRWN 28 Drawing Page(s)

LN.CNT 3127

L5 ANSWER 4 OF 12 USPATFULL

AB The invention relates to the use of pharmacologically valuable pyridyl alkane, pyridyl alkene and/or pyridyl alkine acid amides according to general formula (I) in the treatment of tumors or for immunosuppression. ##STR1##

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AN 2002:239033 USPATFULL

TI Use of pyridyl alkane, pyridyl alkene and/or pyridyl alkine acid amides in the treatment of tumors or for immunosuppression

IN Biedermann, Elfi, Vaterstetten, GERMANY, FEDERAL REPUBLIC OF Hasmann, Max, Neuried, GERMANY, FEDERAL REPUBLIC OF Loser, Roland, Feldafing, GERMANY, FEDERAL REPUBLIC OF Rattel, Benno, Munich, GERMANY, FEDERAL REPUBLIC OF Reiter, Friedemann, Putzbrunn, GERMANY, FEDERAL REPUBLIC OF Schein, Barbara, Neufahrn, GERMANY, FEDERAL REPUBLIC OF Seibel, Klaus, Grafelfing, GERMANY, FEDERAL REPUBLIC OF

PA Klinge Pharma GmbH, Munich, GERMANY, FEDERAL REPUBLIC OF (non-U.S. corporation)

PI US 6451816

B1 20020917

Vogt, Klaus, Munich, GERMANY, FEDERAL REPUBLIC OF

AI US 1998-216482

19981218 (9)

RLI Continuation of Ser. No. WO 1997-EP3244, filed on 20 Jun 1997

DT Utility

FS GRANTED

EXNAM Primary Examiner: Rotman, Alan L.; Assistant Examiner: Desai, Rita

LREP Fitch, Even, Tabin, & Flannery

CLMN Number of Claims: 18

ECL Exemplary Claim: 1

DRWN 0 Drawing Figure(s); 0 Drawing Page(s)

LN.CNT 4285

L5 ANSWER 5 OF 12 USPATFULL

The invention relates to new pyridyl alkane acid amides according to general formula (I) as well as methods for their production, medicaments containing these compounds as well as their medical use, especially in the treatment of tumors or for immunosuppression. ##STR1##

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AN 2002:224728 USPATFULL

TI Pyridyl alkane acid amides as cytostatics and immunosuppressives

IN Biedermann, Elfi, Vaterstetten, GERMANY, FEDERAL REPUBLIC OF

Hasmann, Max, Neuried, GERMANY, FEDERAL REPUBLIC OF

 Loser, Roland, Feldafing, GERMANY, FEDERAL REPUBLIC OF Rattel, Benno, Munich, GERMANY, FEDERAL REPUBLIC OF Reiter, Friedemann, Putzbrunn, GERMANY, FEDERAL REPUBLIC OF Schein, Barbara, Neufahrn, GERMANY, FEDERAL REPUBLIC OF

Seibel, Klaus, Gra felfing, GERMANY, FEDERAL REPUBLIC OF

Vogt, Klaus, Munich, GERMANY, FEDERAL REPUBLIC OF

PA Klinge Pharma GmbH, Munich, GERMANY, FEDERAL REPUBLIC OF (non-U.S. corporation)

PI US 6444823

B1 20020903

AI US 1998-216075

19981218 (9)

RLI Continuation of Ser. No. WO 1997-EP3243, filed on 20 Jun 1997

PRAI DE 1996-DE19624704 19960620

DT Utility

FS GRANTED

EXNAM Primary Examiner: Rotman, Alan L.; Assistant Examiner: Desai, Rita

LREP Fitch, Even, Tabin & Flannery

CLMN Number of Claims: 15

ECL Exemplary Claim: 1

DRWN 0 Drawing Figure(s); 0 Drawing Page(s)

LN.CNT 3772

L5 ANSWER 6 OF 12 USPATFULL

The invention relates to a stabilized medicament with an amount of active ingredients containing cysteine groups and NSAID compounds, wherein a stabilization of the combination, especially the active ingredients containing the cysteine group, can be conducted with a mixture of at least three anti-oxidative components. The therapeutic and prophylactic use of this medicament stabilized in this manner lies in the field of the prevention and therapy of inflammatory diseases among the fields of medical indications.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AN 2002:67202 USPATFULL

TI Stabilized medicament containing cysteinyl derivatives

IN Stanislaus, Fritz, Muenchen, GERMANY, FEDERAL REPUBLIC OF

PI US 2002037855 A1 20020328

AI US 2001-816769 A1 20010322 (9)

RLI Continuation of Ser. No. US 2000-403160, filed on 5 May 2000, ABANDONED A 371 of International Ser. No. WO 1997-EP1941, filed on 18 Apr 1997, UNKNOWN

DT Utility

FS APPLICATION

LREP HELLER EHRMAN WHITE & MCAULIFFE LLP, 275 MIDDLEFIELD ROAD, MENLO PARK, CA, 94025-3506

CLMN Number of Claims: 16

ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 1064
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 7 OF 12 USPATFULL

The invention relates to aqueous dispersions of sparingly water-soluble or water-insoluble organic UV filter substances, which comprise at least one sparingly water-soluble or water-insoluble organic UV filter substance as colloidally disperse phase in amorphous or partially amorphous form.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AN 2001:160697 USPATFULL

TI Aqueous dispersion of water-insoluble organic UV filter substances

IN Heger, Robert, Heidelberg, Germany, Federal Republic of Auweter, Helmut, Limburgerhof, Germany, Federal Republic of Dausch, Wilma M., Limburgerhof, Germany, Federal Republic of Zwissler, Georg Konrad, Heidelberg, Germany, Federal Republic of Wunsch, Thomas, Speyer, Germany, Federal Republic of

PI US 2001022965 A1 20010920 US 6531117 B2 20030311

AI US 2001-771594 A1 20010130 (9)

PRAI DE 2000-10007116 20000217 DE 2000-10042444 20000829

DT Utility FS APPLICATION

LREP Messrs. Keil & Weinkauf, 1101 Connecticut Ave., N.W., Washington, DC, 20036

CLMN Number of Claims: 34 ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 1586

L5 ANSWER 8 OF 12 USPATFULL

The invention relates to medicament excipient particles which are suitable for tissue-specific application of a medicament, especially to the central nervous system (CNS). The invention particles can be loaded with or be free pf the active substance. At least one detection protein is bonded to the particle surface or alternatively, the particle surface is modified in such a way that a detection protein bonds with it on contact.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AN 2001:152938 USPATFULL

TI Medicament excipient particles for tissue-specific application of a medicament

IN Muller, Ranier H., Berlin, Germany, Federal Republic of Luck, Martin, Berlin, Germany, Federal Republic of Kreuter, Jorg, Bad Homburg, Germany, Federal Republic of

PA DSS Drug Delivery Service Gesellschaft zur Forderung der Foshung In Phamazeutischer Technologi und Biopharmazie mbH, Kronshagen, Germany, Federal Republic of (non-U.S. corporation)

PI US 6288040 B1 20010911

WO 9920256 19990429

AI US 2000-529600 20000621 (9)

WO 1998-EP6429 19981013

20000621 PCT 371 date 20000621 PCT 102(e) date

PRAI DE 1997-19745950 19971017

DT Utility FS GRANTED

EXNAM Primary Examiner: Davenport, Avis M.

LREP Melcher, Jeffrey S.Manelli Denison & Selter, PLLC

CLMN Number of Claims: 55 ECL Exemplary Claim: 1

DRWN 4 Drawing Figure(s); 4 Drawing Page(s)

LN.CNT 999

L5

PATENT APPLICATION - PATENTANMELDUNG - DEMANDE DE BREVET

1031564 EUROPATFULL ED 20000910 EW 200035 FS OS ΑN Inhibitors of cellular nicotinamide mononucleotide formation and their TIEN use in cancer therapy. Hemmer der Nicotinamidmonononukleotide-Bildung und deren Verwendung zur TIDE Krebstherapie. Inhibiteurs de la formation du nicotinamide mononucleotide et leur TIFR utilisation dans le traitement du cancer. Biedermann, Elfi, Zugspitzstrasse 93, 85591 Vaterstetten, DE; ΙN Eisenburger, Rolf Dr., Rathausstrasse 4, 85614 Kirchseeon, DE; Hasmann, Max Dr., Lerchenweg 9, 82061 Neuried, DE; Loeser, Roland Dr., Fichtenweg 2, 82340 Feldafing, DE; Rattel, Benno Dr., Eichelhaeherstrasse 3, 81249 Muenchen, DE; Reiter, Friedemann Dr., Zugspitzstrasse 36, 85640 Putzbrunn, DE; Schein, Barbara, Sudetenweg 3, 85375 Neufahrn, DE; Schemainda, Isabel, Hoerwarthstrasse 47, 80804 Muenchen, DE; Schulz, Michael Dr., Sonnenstrasse 6, 85609 Aschheim, DE; Seibel, Klaus Prof. Dr., Haberlstrasse 9, 82166 Graefelfing, DE; Vogt, Klaus Dr., St.-Cajetan-Strasse 32, 81669 Muenchen, DE; Wosikowski, Katja Dr., Seerosenstrasse 3, 85586 Poing, DE Klinge Pharma GmbH, Berg-am-Laim-Strasse 129, D-81673 Muenchen, DE PA PAN 283200 HOFFMANN - EITLE, Patent- und Rechtsanwaelte Arabellastrasse 4, 81925 AG Muenchen, DE AGN 101511 BEPA2000067 EP 1031564 A1 0024 OS SO Wila-EPZ-2000-H35-T1a DTPatent LΑ Anmeldung in Englisch; Veroeffentlichung in Englisch R AT; R BE; R CH; R CY; R DE; R DK; R ES; R FI; R FR; R GB; R GR; R IE; DS R IT; R LI; R LU; R MC; R NL; R PT; R SE; R AL; R LT; R LV; R MK; R RO;

PIT EPA1 EUROPAEISCHE PATENTANMELDUNG PI EP 1031564 A1 20000830 OD 20000830

AI EP 1999-103814 19990226

GRANTED PATENT - ERTEILTES PATENT - BREVET DELIVRE

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934309 EUROPATFULL ED 20020917 EW 200237 FS PS
AN
       NEW PYRIDYL ALKANE ACID AMIDES AS CYTOSTATICS AND IMMUNOSUPPRESSIVES.
TIEN
       PYRIDYLALKAN-SAEUREAMIDE ALS CYTOSTATIKA UND IMMUNOSUPRESSIVE
TIDE
       ARZNEIMITTEL.
       NOUVEAUX AMIDES A ACIDES PYRIDYL-ALCANE UTILISES COMME CYTOSTATIQUES ET
TIFR
       IMMUNOSUPPRESSEURS.
       BIEDERMANN, Elfi, Zugspitzstrasse 93, D-85591 Vaterstetten, DE;
IN
       HASMANN, Max, Lerchenweg 9, D-82061 Neuried, DE;
       LOeSER, Roland, Fichtenweg 2, D-82340 Feldafing, DE;
       RATTEL, Benno, Eichelhaeherstrasse 3, D-81249 Munich, DE;
       REITER, Friedemann, Zugspitzstrasse 36, D-85640 Putzbrunn, DE;
       SCHEIN, Barbara, Sudetenweg 4, D-85375 Neufahrn, DE;
       SEIBEL, Klaus, Haberlstrasse 9, D-82166 Graefelfing, DE;
       VOGT, Klaus, Balanstrasse 63, D-81541 Munich, DE
       Fujisawa Deutschland GmbH, Berg-am-Laim-Strasse 129, 81673 Muenchen, DE
PA
PAN
       HOFFMANN - EITLE, Patent- und Rechtsanwaelte Arabellastrasse 4, 81925
AG
       Muenchen, DE
AGN
       101511
       BEPB2002065 EP 0934309 B1 0123
OS
       Wila-EPS-2002-H37-T1
SO
       Patent
DT
       Anmeldung in Englisch; Veroeffentlichung in Englisch
LΑ
       R AT; R BE; R CH; R DE; R DK; R ES; R FI; R FR; R GB; R GR; R IE; R IT;
DS
       R LI; R LU; R MC; R NL; R PT; R SE
PIT
       EPB1 EUROPAEISCHE PATENTSCHRIFT
                                          (Internationale Anmeldung)
PΙ
       EP 934309
                            B1 20020911
OD
                               19990811
       EP 1997-929240
                               19970620
ΑI
                               19960620
       DE 1996-19624704
PRAI
       WO 97-EP3243
                          970620 INTAKZ
RLI
                          971224 INTPNR
       WO 9748695
       EP 330026 A
                               EP 343307
REP
                               WO 91-15485 A
       WO 91-15484 A
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GRANTED PATENT - ERTEILTES PATENT - BREVET DELIVRE

923570 EUROPATFULL ED 20021007 EW 200239 FS PS PYRIDYL ALKENE- AND PYRIDYL ALKINE- ACID AMIDES AS CYTOSTATICS AND TIEN IMMUNOSUPPRESSIVES. PYRIDYLALKEN- UND PYRIDYLALKIN-SAEUREAMIDE ALS CYTOSTATIKA UND TIDE IMMUNOSUPPRESSIVE ARZNEIMITTEL. AMIDES PYRIDYL-ALCENE ET PYRIDYL-ALCYNE ACIDES UTILISES COMME TTFR · CYTOSTATIQUES ET IMMUNOSUPPRESSEURS. BIEDERMANN, Elfi, Zugspitzstrasse 93, D-85591 Vaterstetten, DE; ΙN HASMANN, Max, Lerchenweg 9, D-82061 Neuried, DE; LOeSER, Roland, Fichtenweg 2, D-82340 Feldafing, DE; RATTEL, Benno, Eichelhaeherstrasse 3, D-81249 Munich, DE; REITER, Friedemann, Zugspitzstrasse 36, D-85640 Putzbrunn, DE; SCHEIN, Barbara, Sudetenweg 4, D-85375 Neufahrn, DE; SEIBEL, Klaus, Haberlstrasse 9, D-82166 Graefelfing, DE; VOGT, Klaus, Balanstrasse 63, D-81541 Munich, DE Fujisawa Deutschland GmbH, Berg-am-Laim-Strasse 129, 81673 Muenchen, DE PA PAN 283202 HOFFMANN - EITLE, Patent- und Rechtsanwaelte Arabellastrasse 4, 81925 ΑG Muenchen, DE AGN 101511 BEPB2002069 EP 0923570 B1 0124 os SO Wila-EPS-2002-H39-T1 Patent DT Anmeldung in Englisch; Veroeffentlichung in Englisch LA R AT; R BE; R CH; R DE; R DK; R ES; R FI; R FR; R GB; R GR; R IE; R IT; DS R LI; R LU; R MC; R NL; R PT; R SE EPB1 EUROPAEISCHE PATENTSCHRIFT (Internationale Anmeldung) PIT EP 923570 B1 20020925 PΙ OD 19990623 19970620 ΑI EP 1997-928261 19960620 PRAI DE 1996-19624659 WO 97-EP3245 970620 INTAKZ RLI WO 9748696 971224 INTPNR EP 330026 EP 343307 REP Α

GRANTED PATENT - ERTEILTES PATENT - BREVET DELIVRE

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912176 EUROPATFULL ED 20021007 EW 200239 FS PS
AN
       USE OF PYRIDYL ALKANE, PYRIDYL ALKENE AND/OR PYRIDYL ALKINE ACID AMIDES
TIEN
       IN THE TREATMENT OF TUMORS OR FOR IMMUNOSUPPRESSION.
       VERWENDUNG VON PYRIDYL-ALKAN-, PYRIDIYL ALKAN- UND/ODER PYRIDYL-SAEUREN
TIDE
       AMIDEN ZUR BEHANDLUNG VON TUMOREN ODER FUER IMMUNSUPPRESSION.
       UTILISATION D'AMIDES PYRIDYL-ALCANE, PYRIDYL-ALCENE ET/OU PYRIDYL-ALCYNE
TIFR
       ACIDES DANS LE TRAITEMENT DES TUMEURS ET POUR L'IMMUNOSUPPRESSION.
       BIEDERMANN, Elfi, Zugspitzstrasse 93, D-85591 Vaterstetten, DE;
ΙN
       HASMANN, Max, Lerchenweg 9, D-82061 Neuried, DE;
       LOeSER, Roland, Fichtenweg 2, D-82340 Feldafing, DE;
       RATTEL, Benno, Eichelhaeherstrasse 3, D-81249 Munich, DE;
       REITER, Friedemann, Zugspitzstrasse 36, D-85640 Putzbrunn, DE;
       SCHEIN, Barbara, Sudetenweg 4, D-85375 Neufahrn, DE;
       SEIBEL, Klaus, Haberlstrasse 9, D-82166 Graefelfing, DE;
       VOGT, Klaus, Balanstrasse 63, D-81541 Munich, DE
       Fujisawa Deutschland GmbH, Berg-am-Laim-Strasse 129, 81673 Muenchen, DE
PA
PAN
       283202
       HOFFMANN - EITLE, Patent- und Rechtsanwaelte Arabellastrasse 4, 81925
ΑG
       Muenchen, DE
AGN
       101511
       BEPB2002068 EP 0912176 B1 0152
OS
SO
       Wila-EPS-2002-H39-T1
DT
       Patent
       Anmeldung in Englisch; Veroeffentlichung in Englisch
LΑ
       R AT; R BE; R CH; R DE; R DK; R ES; R FI; R FR; R GB; R GR; R IE; R IT;
DS
       R LI; R LU; R MC; R NL; R PT; R SE
       EPB1 EUROPAEISCHE PATENTSCHRIFT
                                         (Internationale Anmeldung)
PIT
       EP 912176
                            B1 20020925
PΙ
                               19990506
OD
ΑI
       EP 1997-928260
                               19970620
PRAI
       DE 1996-19624668
                               19960620
       WO 97-EP3244
                          970620 INTAKZ
RLI
                          971224 INTPNR
       WO 9748397
REP
       EP 210782 A
                               EP 330026
       EP 343307
                 Α
                               WO 91-15484 A
       WO 91-15485 A
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=> log y COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	71.48	71.69
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-0.65	-0.65

STN INTERNATIONAL LOGOFF AT 15:29:15 ON 03 APR 2003